



Department of Vermont Health Access
NOB 1 South, 280 State Drive
Waterbury, Vermont 05671-1010

~ HUB (OTP) BUPRENORPHINE Prior Authorization Form ~

All requests for Suboxone® Film > 24mg, Buprenorphine/Naloxone tablets > 24mg, and Buprenorphine monotherapy must be reviewed by the Change Healthcare Clinical Call Center. Documentation must accompany this form. For questions, please contact the Change Healthcare help desk at 1-844-679-5363.

Submit request via Fax: 844-679-5366

Prescribing physician:

Name: _____

NPI: _____

Specialty: _____

Phone#: _____

Fax#: _____

Address: _____

Contact Person at HUB (OTP): _____

Member:

Name: _____

Medicaid ID#: _____

Date of Birth: _____ Sex: _____

Diagnosis: _____

Date of Admission to HUB: _____

CHECK HERE IF PATIENT IS ADAP UNINSURED ☐

Request is from the following HUB location: _____ / _____

Name

NPI

☐ **Suboxone® Film > 24 mg** Dose per day requested: _____mg

* Clinical note/letter from prescriber that documents the prescriber's clinical rationale for requesting Suboxone® Film >24mg must be attached (REQUIRED). Requests for doses >24mg will require review by DVHA Medical Director.

☐ **Buprenorphine/Naloxone tablets > 24 mg** Dose per day requested: _____mg

* Clinical note/letter from prescriber that documents the prescriber's clinical rationale for requesting buprenorphine/naloxone tablets >24mg must be attached (REQUIRED). Requests for doses >24mg will require review by DVHA Medical Director.

☐ **Buprenorphine tablets (mono formulation, any dose)** Dose per day requested: _____mg

☐ Pregnancy DUE DATE: _____ ☐ Pregnancy test/ultrasound result/lab attached (REQUIRED)

☐ Breastfeeding an opiate dependent baby (baby is being administered morphine or methadone for opiate withdrawal symptoms)

*Clinical note/letter from a pediatrician/neonatologist must be attached (REQUIRED)

☐ Using buprenorphine mono to switch from methadone to Suboxone®
Dates buprenorphine mono will be administered: _____

☐ Using buprenorphine mono due to provider observed reaction to both Suboxone® Film and Buprenorphine/Naloxone tablets severe enough to require discontinuation

* Clinical documentation is submitted detailing a provider-observed reaction to both Suboxone films and buprenorphine/naloxone tablets severe enough to require discontinuation (documentation of measures tried to mitigate/manage symptoms is required).

Prescriber Signature: _____ (stamps not acceptable) Date of request: _____

